AMENDMENTS TO THE CLAIMS

This listing of claims will replace all prior versions and listings of claims in the application:

LISTING OF CLAIMS:

- 1. (original): A process for preparing crystalline parahydroxybenzoic acid anhydride, comprising the step of precipitating and isolating parahydroxybenzoic acid in an aqueous solvent at a temperature equal to or above the transition temperature of parahydroxybenzoic acid.
- 2. (currently amended): The process for preparing crystalline parahydroxybenzoic acid anhydride according to claim 1, wherein the precipitating and isolating step is performed at a temperature which is in the range from the transition temperature to said the transition temperature + 30°C.
- 3. (original): A process for preparing crystalline parahydroxybenzoic acid anhydride, comprising the step of precipitating and isolating parahydroxybenzoic acid with acid from a solution of parahydroxybenzoate in an aqueous solvent at a temperature equal to or above the transition temperature of parahydroxybenzoic acid.
- 4. (original): A process for preparing crystalline parahydroxybenzoic acid anhydride, comprising the steps of: precipitating parahydroxybenzoic acid in an aqueous solvent with acid, heating the parahydroxybenzoic acid precipitates to dissolve the same, and reprecipitating and isolating the parahydroxybenzoic acid at a temperature equal to or above the transition temperature of parahydroxybenzoic acid.

5. (currently amended): A process for preparing crystalline parahydroxybenzoic acid anhydride, comprising the steps of:

providing a suspension of parahydroxybenzoic acid monohydride in an aqueous solvent; preparing a solution of parahydroxybenzoic acid in an aqueous solvent by heating the suspension; and

precipitating crystalline parahydroxybenzoic acid anhydride by keeping said solution at a temperature equal to or above the transition temperature of parahydroxybenzoic acid; and isolating the crystalline parahydroxybenzoic acid anhydride at a temperature equal to or above the transition temperature of parahydroxybenzoic acid.

6. (currently amended): A process for preparing crystalline parahydroxybenzoic acid anhydride, comprising the steps of:

<u>providing preparing</u>-a suspension of parahydroxybenzoic acid<u>monohydride</u> in an aqueous solvent,

<u>changing parahydroxybenzoic acid monohydride to parahydroxybenzoic acid anhydride</u>

<u>by heating the suspension to a temperature equal to or above the transition temperature of parahydroxybenzoic acid, and</u>

isolating the crystalline parahydroxybenzoic acid anhydride at a temperature equal to or above the transition temperature of parahydroxybenzoic acid.

7. (currently amended): The process for preparing crystalline parahydroxybenzoic acid anhydride according to any one of claims 1 to 6 claim 1, 2, 3, 4, 5 or 6, wherein the aqueous solvent is water and the transition temperature of parahydroxybenzoic acid is 52 to 54°C.

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8. (currently amended): Crystalline parahydroxybenzoic acid anhydride, wherein the specific surface area of particles that can pass through a 100 mesh (150 μ m) sieve and can not pass through a 140 mesh (106 μ m) sieve is equal to or less than 0.3 m²/g.

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- 9. (original): The crystalline parahydroxybenzoic acid anhydride according to claim 8, wherein the angle of repose is equal to or less than 45°.
- 10. (original): The crystalline parahydroxybenzoic acid anhydride according to claim 8 or 9, wherein the compression ratio calculated according to the following formula is equal to or less than 10%: (packed bulk density-aerated bulk density)/packed bulk density x 100.